CLAIMS

1. A triaza-cyclopenta[cd]indene derivative represented by the following formula [I]:

$$R^{1}$$
 N
 R^{5}
 R^{3}
 N
 R^{4}
 R^{5}

(wherein R¹ and R² are the same or different, and independently are hydrogen, C₁. 6alkyl, C₃₋₇cycloalkyl, C₃₋₇cycloalkyl-C₁₋₆alkyl, C₁₋₆alkyl, C₁₋₆alkyl, hydroxy-C₁. 6alkyl, cyano-C₁₋₆alkyl, carbamoyl-C₁₋₆alkyl or di(C₁₋₆alkyl)amino-C₁₋₆alkyl, cyano, carbamoyl or aryl;

R³ is hydrogen, C₁₋₆alkyl, C₃₋₇cycloalkyl, C₃₋₇cycloalkyl-C₁₋₆alkyl, halogen, C₁₋₆alkoxy, C₃₋₇cycloalkyloxy, C₁₋₆alkylthio or -N(R⁶)R⁷;

R⁴ is hydrogen, C₁₋₆alkyl, C₃₋₇cycloalkyl or C₃₋₇cycloalkyl-C₁₋₆alkyl; R⁵ is hydrogen, C₁₋₆alkyl, aryl-C₁₋₆alkyl or carbamoyl;

Ar is aryl or heteroaryl which aryl or heteroaryl is unsubstituted or substituted with 1 or more substituents, which are the same or different, selected from the group consisting of halogen, C₁₋₆alkyl, C₃₋₇cycloalkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₁₋₆alkoxy, C₁₋₆alkylthio, C₁₋₆alkylsulfinyl, C₁₋₆alkylsulfonyl, cyano, nitro, hydroxy, -CO₂R⁸, -C(=O)R⁹, -CONR¹⁰R¹¹, -OC(=O)R¹², -NR¹³CO₂R¹⁴, -S(=O)₁NR¹⁵R¹⁶, trifluoromethyl, trifluoromethoxy, difluoromethoxy, fluoromethoxy and -N(R¹⁷)R¹⁸;

R⁸ and R¹⁴ are the same or different, and independently are hydrogen or C₁₋₅alkyl, C₃₋₈cycloalkyl, C₃₋₈cycloalkyl-C₁₋₅alkyl, aryl or aryl-C₁₋₅alkyl;

R⁶, R⁷, R⁹, R¹⁰, R¹¹, R¹², R¹³, R¹⁵, R¹⁶, R¹⁷ and R¹⁸ are the same or different, and independently are hydrogen, C₁₋₆alkyl or C₃₋₇cycloalkyl;

r is 1 or 2) or individual isomers thereof or racemic or non-racemic mixtures of isomers thereof, or pharmaceutically acceptable salts and hydrates thereof.

2. The triaza-cyclopenta[cd]indene derivative according to claim 1 represented by the formula [I], wherein R³ is C₁₋₆alkyl; R⁴ is hydrogen or C₁₋₆alkyl;

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 R^5 is hydrogen or $C_{1\text{-}6}$ alkyl; Ar is phenyl which phenyl is substituted with two or three substituents, which are the same or different, selected from the group consisting of halogen, $C_{1\text{-}3}$ alkyl, $C_{1\text{-}3}$ alkoxy, $C_{1\text{-}3}$ alkylthio, trifluoromethyl, trifluoromethoxy and $-N(R^{17})R^{18}$ (wherein R^{17} and R^{18} are the same or different, and independently are hydrogen or $C_{1\text{-}3}$ alkyl); R^1 , R^2 and R^4 are as defined in claim 1, or pharmaceutically acceptable salts and hydrates thereof.

- 3. The triaza-cyclopenta[cd]indene derivative according to claim 1 represented by the formula [I], wherein R^3 is C_{1-3} alkyl; R^5 is hydrogen or C_{1-3} alkyl; Ar is phenyl which phenyl is substituted with two or three substituents, which are the same or different, selected from the group consisting of halogen and C_{1-3} alkyl; R^1 , R^2 and R^4 are as defined in claim 1, or pharmaceutically acceptable salts and hydrates thereof.
- 4. An antagonist for CRF receptors, comprising a triaza-cyclopenta[cd]indene derivative, a pharmaceutically acceptable salt thereof or its hydrate according to any one of claims 1 to 3, as an active ingredient.
- 5. Use of a triaza-cyclopenta[cd]indene derivative, a pharmaceutically acceptable salt thereof or its hydrate according to any one of claim 1 to 3, for the manufacture of a therapeutic agent as an antagonist for CRF receptors.